

## **Refine Search**

#### Search Results -

Terms	Documents					
L6 with L4	14					

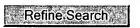
US Pre-Grant Publication Full-Text Database US Patents Full-Text Database

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### **Search History**

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DB=P	GPB,USPT; PLUR=YES; OP=AND		
<u>L7</u>	L6 with 14	14	<u>L7</u>
<u>L6</u>	11 with 12	3654	<u>L6</u>
<u>L5</u>	L4 and 13	286	<u>L5</u>
<u>L4</u>	fabry adj disease	818	<u>L4</u>
<u>L3</u>	11 and L2	6208	<u>L3</u>
<u>L2</u>	gene adj therapy	34350	<u>L2</u>
L1	(enzyme or protein or polypeptide) near3 therapy	9253	L1

**END OF SEARCH HISTORY** 

# Generate Collection Print

## Search Results - Record(s) 1 through 14 of 14 returned.

☐ 1. 20040204379. 16 Jan 04. 14 Oct 04. Combination enzyme replacement, gene therapy and small molecule therapy for lysosomal storage diseases. Cheng, Seng H., et al. 514/44; A61K048/00.
☐ 2. <u>20040029779</u> . 04 Apr 03. 12 Feb 04. Methods of enhancing lysosomal storage disease therapy by modulation of cell surface receptor density. Zhu, Yunxiang, et al. 514/3; 424/85.1 424/85.2 514/12 514/179 514/23 514/573 A61K038/28 A61K038/19 A61K038/20 A61K031/70 A61K031/573 A61K031/557.
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L6 with L4	14

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- Douglas A.; Williams, Melanie D.; Schuetz, Thomas J.; Daniel, Peter F.
- TI Purification of recombinant  $\alpha$ -galactosidase A and its glycosylation

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- L8 ANSWER 4 OF 12 CAPLUS COPYRIGHT 2006 ACS on STN
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- TI Production in yeast of  $\alpha$ -galactosidase A, a lysosomal enzyme applicable to enzyme replacement therapy for Fabry disease
- SO Glycobiology (2002), 12(12), 821-828 CODEN: GLYCE3; ISSN: 0959-6658
- L8 ANSWER 5 OF 12 CAPLUS COPYRIGHT 2006 ACS on STN
- AU Branton, Mary H.; Schiffmann, Raphael; Sabnis, Sharda G.; Murray, Gary J.; Quirk, Jane M.; Altarescu, Gheona; Goldfarb, Lev; Brady, Roscoe O.; Balow, James E.; Austin, Howard A., III; Kopp, Jeffrey B.
- TI Natural history of Fabry renal disease: Influence of  $\alpha$ -galactosidase A activity and genetic mutations on clinical course
- SO Medicine (Baltimore, MD, United States) (2002), 81(2), 122-138 CODEN: MEDIAV; ISSN: 0025-7974
- L8 ANSWER 6 OF 12 CAPLUS COPYRIGHT 2006 ACS on STN
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- Selden, Richard F.; Borowski, Marianne; Gillispie, Frances P.; Kinoshita, IN Carol M.; Treco, Douglas A.; Williams, Melanie D. Gene and enzyme replacement therapy for
- ΤI alpha.-galactosidase A deficiency
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	US 6458574	B1	20021001	US 1999-266014	19990311
	US 6566099	B1	20030520	US 2000-491759	20000127
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- L8 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2006 ACS on STN
- IN Selden, Richard F.; Borowski, Marianne; Gillespie, Frances P.; Kinoshita, Carol M.; Treco, Douglas A.; Williams, Melanie D.
- Gene and enzyme replacement therapy for . alpha. -galactosidase A deficiency
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ΡI	WO	9811	206			A2		1998	0319	1	WO 1	997-1	US16	603		1:	9970	912
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- TI Structural organization and expression of the mouse gene encoding .alpha.-galactosidase A
- SO Gene (1995), 166(2), 277-80 CODEN: GENED6; ISSN: 0378-1119
- L8 ANSWER 10 OF 12 CAPLUS COPYRIGHT 2006 ACS on STN
- IN Desnick, Robert J.; Bishop, David F.; Ioannou, Yiannis A.
- TI Biologically active human  $\alpha$ -galactosidase A and its manufacture by expression of the cloned gene
- SO PCT Int. Appl., 154 pp.

CODEN: PIXXD2

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- L8 ANSWER 11 OF 12 CAPLUS COPYRIGHT 2006 ACS on STN
- IN Calhoun, David H.; Coppola, George
- TI Recombinant manufacture of human  $\alpha$ -galactosidase for treatment of Fabry's disease
- SO PCT Int. Appl., 43 pp.

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ΡI	WO 9011353	A1 19901004	WO 1990-US1571	19900323
	RW: AT, BE, CH,	DE, DK, ES, FR,	GB, IT, LU, NL, SE	
	EP 463109	A1 19920102	EP 1990-905932	19900323
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- TI Structural organization of the human .alpha.galactosidase A gene: further evidence for the absence of a 3' untranslated region
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- AU Grabowski, Gregory A.; Hopkin, Robert J.
- TI Enzyme therapy for lysosomal storage disease: Principles, practice, and prospects
- SO Annual Review of Genomics and Human Genetics (2003), 4, 403-436 CODEN: ARGHC4; ISSN: 1527-8204
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- AU Chiba, Yasunori
- TI The development of therapeutic methods of lysosomal diseases
- SO Igaku no Ayumi (2003), 207(5), 387-392 CODEN: IGAYAY; ISSN: 0039-2359
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- AU Mohrenschlager Matthias; Braun-Falco Markus; Ring Johannes; Abeck Dietrich
- TI Fabry disease: recognition and management of cutaneous manifestations.
- SO American journal of clinical dermatology, (2003) Vol. 4, No. 3, pp. 189-96. Ref: 54

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- AU Blom Daniel; Speijer Dave; Linthorst Gabor E; Donker-Koopman Wilma G; Strijland Anneke; Aerts Johannes M F G
- TI Recombinant enzyme therapy for Fabry disease: absence of editing of human alpha-galactosidase A mRNA.
- SO American journal of human genetics, (2003 Jan) Vol. 72, No. 1, pp. 23-31. Electronic Publication: 2002-12-06.
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- AU Brady R O
- TI Gaucher and Fabry diseases: from understanding pathophysiology to rational therapies.
- SO Acta paediatrica (Oslo, Norway : 1992). Supplement, (2003 Dec) Vol. 92, No. 443, pp. 19-24.
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- L13 ANSWER 25 OF 49 CAPLUS COPYRIGHT 2006 ACS on STN
- IN Davidson, Beverly L.; Mao, Qinwen; Xia, Haibin
- TI Gene therapy vectors comprising lysosomal enzyme and modified Tat protein transduction domain for treating genetic diseases and cancer
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